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## ORIGINAL ARTICLE

# Effect of alirocumab on lipids and lipoproteins in individuals with metabolic syndrome without diabetes: Pooled data from 10 phase 3 trials

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**Aims:** This analysis assessed the efficacy and safety of alirocumab, a proprotein convertase subtilisin/kexin type 9 inhibitor, in patients with or without metabolic syndrome (MetS) using pooled data from 10 phase 3 ODYSSEY trials.

**Materials and Methods:** Data from 4983 randomized patients (1940 with MetS; 1642 with diabetes excluded) were assessed in subgroups by MetS status. Efficacy data were analysed in 4 pools per study design: 2 placebo-controlled pools (1 using alirocumab 150 mg every 2 weeks [Q2W], 1 using 75/150 mg Q2W) with background statin, and 2 ezetimibe-controlled pools (both alirocumab 75/150 mg Q2W), 1 with and 1 without background statin. Alirocumab 75/150 mg indicates possible dose increase from 75 to 150 mg at Week 12 based on Week 8 LDL-C.

**Results:** LDL-C percentage reduction from baseline at Week 24 with alirocumab was 63.9% (MetS) and 56.8% (non-MetS) in the pool of alirocumab 150 mg Q2W, and 42.2% to 52.2% (MetS) and 45.0% to 52.6% (non-MetS) in 3 pools using 75/150 mg Q2W. Levels of other lipid and lipoprotein parameters were also improved with alirocumab treatment, including apolipoprotein B, non-high-density lipoprotein cholesterol (non-HDL-C), lipoprotein(a) and HDL-C. Overall, the percentage change at Week 24 in LDL-C and other lipids and lipoproteins did not vary by MetS status. Adverse event rates were generally similar between treatment groups, regardless of MetS status; injection-site reactions occurred more frequently in alirocumab vs control groups.

**Conclusions:** Across study pools, alirocumab-associated reductions in LDL-C, apolipoprotein B, and non-HDL-C were significant vs control, and did not vary by MetS status.

## KEYWORDS

cardiovascular disease, clinical trial, dyslipidaemia, lipid-lowering therapy

## 1 | INTRODUCTION

Metabolic syndrome (MetS) is defined by a collection of related metabolic and physiological abnormalities, including central obesity, raised serum triglycerides (TGs), reduced high-density lipoprotein cholesterol (HDL-C), glucose intolerance and hypertension.<sup>1–3</sup> MetS has been defined as the presence of 3 or more of the following: elevated

waist circumference (which is assumed in the guidelines if body mass index [BMI] is  $>30 \text{ kg/m}^2$ ); TGs  $\geq 150 \text{ mg/dL}$  or use of TG-lowering medication; HDL-C  $<40 \text{ mg/dL}$  in men or  $<50 \text{ mg/dL}$  in women; blood pressure  $\geq 130/85 \text{ mm Hg}$  or diagnosis of hypertension; and fasting plasma glucose (FPG)  $\geq 100 \text{ mg/dL}$ .<sup>1</sup> MetS is a common syndrome with a rising prevalence worldwide, ranging from approximately 10% to 80% depending on regional variation and population

demographics.<sup>1,2</sup> MetS is associated with an increased risk of cardiovascular (CV) disease (2-fold), type 2 diabetes mellitus (5-fold) and all-cause mortality (1.5-fold).<sup>1,4,5</sup> However, multivariate analyses have shown that components typically associated with MetS (blood pressure, HDL-C and blood glucose), but not MetS itself, are predictors of prevalent coronary heart disease.<sup>6</sup>

While statins are recommended as first-line therapy for reducing levels of LDL-C, not all patients achieve LDL-C lowering with statin therapy sufficient to optimally reduce their CV risk. Additionally, statin intolerance can limit dosage and potency of the statin used, which is a significant factor in the reduced efficacy of statin therapy for some patients.<sup>7</sup> In particular, patients with MetS may not achieve non-HDL-C goals following treatment with statins and other lipid-lowering therapies (LLTs)<sup>8</sup> due to elevated TG levels, which are indicators of very-low-density lipoprotein and remnant cholesterol levels.<sup>9</sup> Individuals with MetS typically exhibit mixed dyslipidaemia, characterised by elevated TGs and lower HDL-C, both of which are often associated with elevated apolipoprotein (Apo) B and non-HDL-C levels. In such cases, non-HDL-C or ApoB may give a better estimate of the concentration of atherogenic particles than low-density lipoprotein cholesterol (LDL-C), and they have been suggested as alternative treatment targets to LDL-C for such individuals.<sup>4</sup>

Alirocumab, a proprotein convertase subtilisin/kexin type 9 inhibitor, has been approved in more than 40 countries, including the USA and the EU, for reducing elevated LDL-C levels.<sup>10,11</sup> Individuals with metabolic syndrome have a range of clinical and metabolic characteristics that may impact the efficacy and/or safety of a PCSK9 inhibitor, including obesity, high triglycerides/low HDL-C, high blood glucose and insulin resistance. For example, PCSK9 appears to have a role in glucose metabolism and PCSK9 levels have been shown to correlate with glycaemic parameters and insulin resistance, although evidence is conflicting among different studies.<sup>12,13</sup> Insulin signalling, which is known to influence LDL-receptor expression (the target of PCSK9), is often dysregulated in metabolic syndrome.<sup>14</sup> However, the use of PCSK9 inhibitors in individuals with metabolic syndrome is not well established. In the present study, we compared the efficacy and safety of alirocumab in patients with hypercholesterolaemia, with or without MetS, using data from 10 studies from the alirocumab ODYSSEY phase

3 clinical trial programme. This analysis contributes to a better understanding of the alirocumab target patient population.

## 2 | METHODS

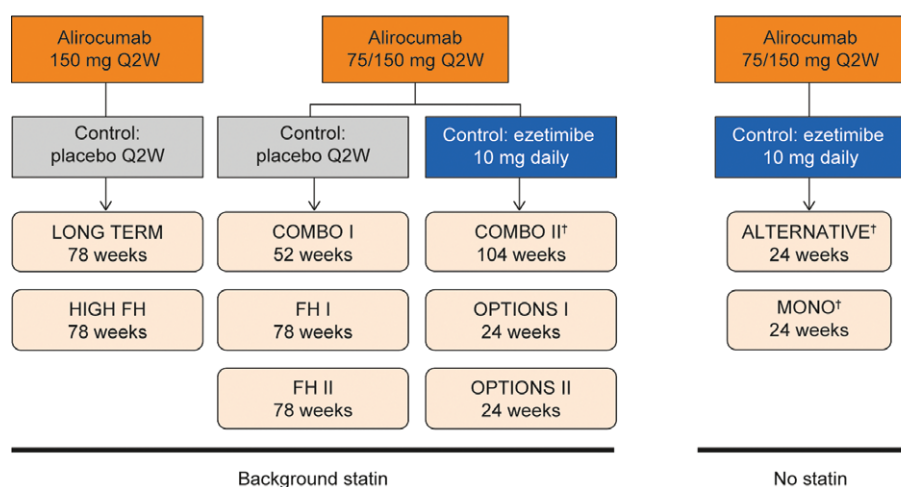
### 2.1 | Study designs and pooling strategy

This analysis includes data from 10 phase 3 ODYSSEY studies (Figure 1), all of which have been described previously.<sup>15–23</sup> Patients ( $n = 4983$ ) were randomized to receive alirocumab or control (placebo or ezetimibe). In 8 of the 10 trials, patients were receiving background statin therapy, with or without other LLTs. The double-blind treatment periods ranged between 24 and 104 weeks. In 8 studies, patients received an initial dose of alirocumab 75 mg every 2 weeks (Q2W), with a possible dose increase to 150 mg Q2W at Week 12, depending on LDL-C levels at Week 8 (alirocumab 75/150 mg Q2W); 2 studies used alirocumab 150 mg Q2W throughout the study. Entry criteria for the ODYSSEY trials included baseline LDL-C  $\geq 70$  mg/dL for those with prior CV events or  $\geq 100$  mg/dL for those without CV events but with other risk factors. Exceptions were the LONG TERM study (LDL-C  $\geq 70$  mg/dL for all patients<sup>19</sup>), the MONO study (LDL-C 100–190 mg/dL for all patients<sup>21</sup>) and the HIGH FH study (LDL-C  $\geq 160$  mg/dL for all patients<sup>23</sup>). All study protocols were approved by the appropriate review boards, and all patients provided written, informed consent.

For the current analysis, efficacy data were analysed in 4 pools according to alirocumab starting dose, control (placebo or ezetimibe) and use of background statin therapy (yes/no), as shown in Figure 1. In the first pool (LONG TERM,<sup>19</sup> HIGH FH<sup>23</sup>), patients received alirocumab 150 mg Q2W vs placebo (with statins); in the second (COMBO I,<sup>18</sup> FH I & II<sup>15</sup>), alirocumab 75/150 mg Q2W vs placebo (with statins); in the third (COMBO II,<sup>17</sup> OPTIONS I & II<sup>16,20</sup>), alirocumab 75/150 mg Q2W vs ezetimibe (with statins); and in the fourth (ALTERNATIVE,<sup>22</sup> MONO<sup>21</sup>), alirocumab 75/150 mg Q2W vs ezetimibe (without statins).

Patient baseline characteristics and safety and efficacy data were further analysed in subgroups, with and without MetS. MetS was defined in this analysis of the ODYSSEY trials using a definition similar to that proposed by the International Atherosclerosis Society,<sup>24</sup> with 2 important differences. Firstly, waist circumference

**FIGURE 1** Overview of ODYSSEY studies included in this analysis. Abbreviations: LLT, lipid-lowering therapy; Q2W, every 2 weeks. †Additional non-statin LLTs were not allowed. Clinicaltrials.gov identifiers: ALTERNATIVE, NCT01709513; COMBO I, NCT01644175; COMBO II, NCT01644188; FH I, NCT01623115; FH II, NCT01709500; HIGH FH, NCT01617655; LONG TERM, NCT01507831; MONO, NCT01644474; OPTIONS I, NCT01730040; OPTIONS II, NCT01730053



**TABLE 1** Patient baseline characteristics and baseline lipids (randomized population)

	ALI 150 vs PBO (with statins)		ALI 75/150 vs PBO (with statins)		ALI 75/150 vs EZE (with statins)		ALI 75/150 vs EZE (without statins)	
	ALI	PBO	ALI	PBO	ALI	EZE	ALI	EZE
Group, n	MetS 599	334	294	137	254	168	75	79
	Non-MetS 429	184	261	131	166	100	60	70
Age, years, mean (SD)	MetS 59.5 (10.5)	60.4 (9.9)	56.7 (12.2)	55.2 (13.1)	61.3 (9.5)	61.9 (9.8)	62.6 (8.2)	63.2 (8.3)
	Non-MetS 58.3 (12.1)	58.6 (12.2)	50.6 (13.6)	52.6 (12.5)	60.7 (9.7)	60.3 (11.2)	61.8 (7.3)	59.9 (10.5)
Male, % (n)	MetS 65.8 (394)	66.2 (221)	60.9 (179)	64.2 (88)	74.0 (188)	70.2 (118)	58.7 (44)	54.4 (43)
	Non-MetS 64.3 (276)	62.5 (115)	52.5 (137)	58.8 (77)	74.1 (123)	70.0 (70)	51.7 (31)	50.0 (35)
Race, white, % (n)	MetS 96.2 (576)	96.4 (322)	94.9 (279)	91.2 (125)	85.4 (217)	91.7 (154)	94.7 (71)	87.3 (69)
	Non-MetS 97.9 (420)	96.7 (178)	93.5 (244)	94.7 (124)	90.4 (150)	88.0 (88)	95.0 (57)	98.6 (69)
BMI, kg/m <sup>2</sup> , mean (SD)	MetS 30.8 (5.3)	30.6 (4.7)	31.0 (4.4)	30.9 (5.8)	31.1 (5.2)	30.7 (5.0)	30.7 (5.1)	30.3 (5.4)
	Non-MetS 26.6 (3.7)	26.5 (3.5)	26.9 (4.0)	27.0 (4.0)	26.3 (3.2)	26.8 (3.5)	25.4 (3.3)	25.1 (3.0)
SBP, mm Hg, mean (SD)	MetS 135.2 (15.7)	135.2 (13.8)	131.5 (13.7)	129.3 (12.5)	132.3 (14.0)	131.9 (11.8)	131.5 (12.6)	130.4 (12.9)
	Non-MetS 124.7 (15.1)	125.5 (15.1)	121.3 (12.9)	122.4 (14.2)	124.7 (14.1)	121.9 (12.9)	123.3 (13.4)	124.4 (13.3)
HbA1c, %, mean (SD)	MetS 5.7 (0.4)	5.6 (0.3)	5.7 (0.4)	5.6 (0.3)	5.7 (0.3)	5.7 (0.4)	5.7 (0.3)	5.6 (0.4)
	Non-MetS 5.6 (0.3)	5.6 (0.3)	5.5 (0.3)	5.5 (0.3)	5.6 (0.3)	5.6 (0.3)	5.5 (0.3)	5.5 (0.3)
FBG, mg/dL, mean (SD)	MetS 101.3 (11.5)	100.0 (11.9)	101.1 (12.3)	100.4 (10.0)	104.2 (11.4)	101.7 (11.8)	101.7 (15.4)	98.6 (10.2)
	Non-MetS 93.6 (9.7)	94.0 (10.8)	92.4 (9.3)	93.3 (12.0)	97.8 (15.2)	96.9 (14.6)	93.9 (11.7)	93.0 (8.5)
Hypertension, % (n)	MetS 80.3 (481)	82.9 (277)	67.3 (198)	65.0 (89)	83.1 (211)	88.1 (148)	72.0 (54)	72.2 (57)
	Non-MetS 37.8 (162)	39.7 (73)	21.8 (57)	22.9 (30)	57.8 (96)	46.0 (46)	20.0 (12)	18.6 (13)
ASCVD, <sup>a</sup> % (n)	MetS 85.8 (514)	88.3 (295)	61.6 (181)	66.4 (91)	91.3 (232)	89.9 (151)	44.0 (33)	43.0 (34)
	Non-MetS 76.5 (328)	77.2 (142)	42.9 (112)	39.7 (52)	90.4 (150)	82.0 (82)	23.3 (14)	15.7 (11)
Very high cardiovascular risk, % (n)	MetS 87.5 (524)	91.3 (305)	63.6 (187)	69.3 (95)	92.5 (235)	90.5 (152)	44.0 (33)	45.6 (36)
	Non-MetS 78.8 (338)	78.3 (144)	44.4 (116)	41.2 (54)	91.0 (151)	82.0 (82)	25.0 (15)	15.7 (11)
LDL-C, mg/dL, mean (SD)	MetS 129.9 (50.6)	128.9 (47.7)	132.6 (48.7)	134.0 (48.6)	111.9 (37.5)	106.1 (38.2)	184.5 (69.3)	174.2 (58.4)
	Non-MetS 131.9 (48.3)	129.4 (46.1)	132.9 (47.0)	131.6 (40.3)	109.7 (38.4)	110.3 (42.8)	177.6 (73.0)	182.7 (75.3)
HDL-C, mg/dL, mean (SD)	MetS 47.7 (11.5)	48.3 (12.2)	47.8 (14.9)	46.4 (13.7)	46.7 (13.1)	47.3 (12.6)	46.9 (13.0)	47.3 (10.1)

TABLE 1 (Continued)

	ALI 150 vs PBO (with statins)		ALI 75/150 vs PBO (with statins)		ALI 75/150 vs EZE (with statins)		ALI 75/150 vs EZE (without statins)	
	ALI	PBO	ALI	PBO	ALI	EZE	ALI	EZE
Non-MetS	55.2 (13.2)	54.9 (11.2)	56.4 (15.6)	55.5 (15.0)	55.0 (12.6)	53.9 (13.5)	60.1 (17.0)	61.9 (18.8)
MetS	145.1 (108.8:198.0)	154.0 (108.8:197.3)	141.0 (97.0:196.0)	134.5 (101.0:174.0)	148.5 (109.0:200.0)	154.5 (112.5:207.0)	174.0 (135.0:235.0)	163.0 (115.0:225.0)
Non-MetS	93.8 (75.0:126.5)	99.1 (73.7:124.9)	93.0 (72.0:121.0)	90.0 (72.0:109.0)	98.0 (76.0:127.0)	96.5 (78.0:118.5)	100.0 (74.5:120.5)	93.0 (77.0:128.0)
MetS	130.5 (96.0:177.9)	145.1 (95.0:196.0)	125.0 (93.5:180.5)	123.0 (97.0:170.0)	131.0 (106.0:208.0)	142.0 (95.0:191.5)	207.0 (132.0:240.0)	197.0 (120.0:232.5)
Non-MetS	88.0 (72.0:111.5)	94.7 (67.7:126.5)	89.0 (70.0:109.0)	90.0 (70.5:104.5)	89.0 (79.5:106.0)	111.0 (107.0:113.0)	122.0 (105.0:125.0)	108.0 (83.0:129.0)
MetS	153.1 (115.9:205.0)	158.0 (115.9:199.0)	170.5 (118.0:211.0)	147.0 (113.0:198.0)	153.0 (110.0:199.0)	159.0 (115.0:218.5)	172.5 (153.0:233.0)	159.0 (112.0:218.0)
Non-MetS	95.6 (75.0:131.0)	99.1 (78.8:124.4)	95.0 (72.0:126.0)	91.0 (73.0:120.0)	99.0 (76.0:129.0)	96.0 (78.0:120.0)	95.0 (74.0:119.0)	91.0 (73.0:128.0)

Abbreviations: ALI, alirocumab; ASCVD, atherosclerotic cardiovascular disease; BMI, body mass index; EZE, ezetimibe; FPG, fasting plasma glucose; HbA1c, glycated haemoglobin; HDL-C, high-density lipoprotein cholesterol; LDL-C, low-density lipoprotein cholesterol; LLTs, lipid-lowering therapies; MetS, metabolic syndrome; PBO, placebo; SBP, systolic blood pressure; SD, standard deviation; TGs, triglycerides.

<sup>a</sup> Including coronary heart disease, peripheral arterial disease and ischaemic stroke; transient ischaemic attack, carotid endarterectomy or carotid artery stent procedure, and renal artery stent procedure were also included in the definition used for the FH II, ALTERNATIVE and OPTIONS I & II trials.

measurements were not performed in the alirocumab phase 3 program; instead, we used BMI as a proxy, as has been done previously.<sup>25</sup> Second, although 1 of the criteria for the International Atherosclerosis Society definition was FPG  $\geq 100$  mg/dL, which would include individuals with type 2 diabetes, we introduced a cut-off of 126 mg/dL for FPG so that individuals with type 2 diabetes would be excluded. This was to allow examination of the specific metabolic syndrome population known to be at risk of developing diabetes and cardiovascular disease. In this analysis, metabolic syndrome was defined as the presence of 3 or more of the following: BMI  $>30$  kg/m<sup>2</sup> for non-Asians or  $>25$  kg/m<sup>2</sup> for Asians; TGs  $\geq 150$  mg/dL or use of TG-lowering medication (which in this analysis included fibrates); HDL-C  $<40$  mg/dL in men or  $<50$  mg/dL in women; blood pressure  $\geq 130/85$  mm Hg or diagnosis of hypertension; and FPG  $\geq 100$  to  $<126$  mg/dL. Patients with type 1 or type 2 diabetes were excluded from the analysis (type 2 diabetes was defined based on medical history, or baseline glycated haemoglobin [HbA1c]  $\geq 6.5\%$ , or 2 FPG values  $\geq 126$  mg/dL at screening and randomization). All other patients (ie, those who did not have MetS or diabetes) were defined as non-MetS for this analysis.

## 2.2 | Endpoints

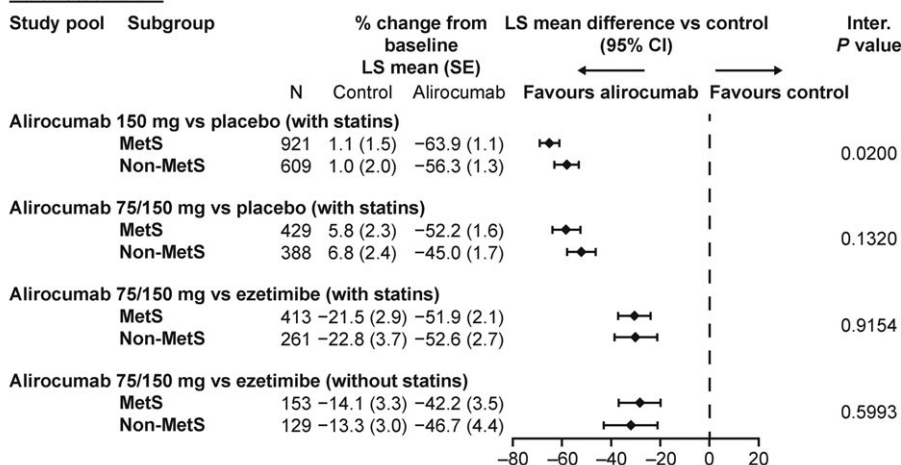
The current pooled analysis uses the same efficacy endpoints as the primary studies. The primary efficacy endpoint in all studies was the percentage change in LDL-C from baseline to Week 24. LDL-C levels were calculated using the Friedewald equation if TGs were  $<400$  mg/dL; in instances above this threshold, LDL-C was determined using the beta-quantification method. Regardless, LDL-C values derived by beta-quantification were not included in the efficacy analysis. Secondary efficacy endpoints included the percentage change from baseline to Week 24 in other lipids, including non-HDL-C, ApoB, lipoprotein (a) (Lp[a]), TGs, HDL-C and TG-rich lipoprotein cholesterol (TRL-C). Non-HDL-C was calculated by subtracting HDL-C from total cholesterol. The concentration of TRL-C was calculated by subtracting HDL-C and calculated LDL-C from total cholesterol, following the method of Nordestgaard et al.<sup>26</sup> Lp(a) and ApoB levels in serum were measured from immunonephelometry by a central laboratory (Medpace Reference Laboratories, Cincinnati, Ohio and Leuven, Belgium, with the exception of the LONG TERM study,<sup>19</sup> which used Covance Central Laboratory, Indianapolis, Indiana). Safety assessments included treatment-emergent adverse events (TEAEs), defined as events occurring from the time of the first dose of study treatment to the last dose, plus 70 days. The change over time in glycaemic parameters (HbA1c and FPG) was also assessed.

## 2.3 | Statistical analysis

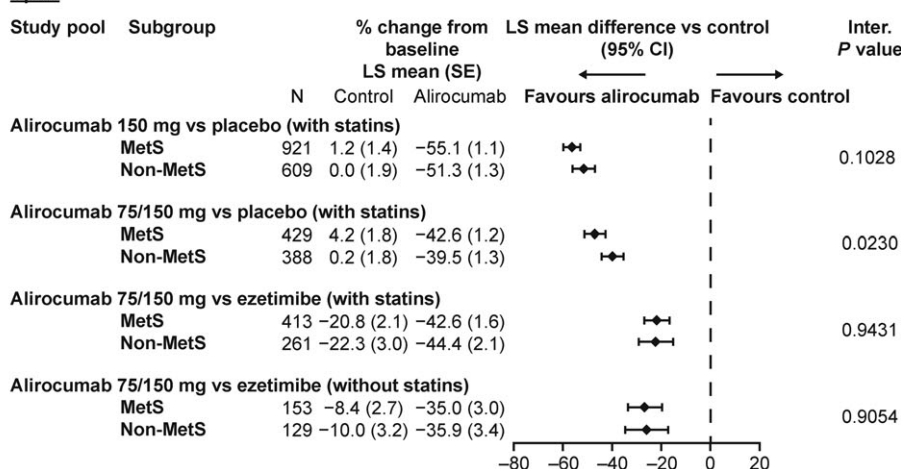
Efficacy data were analysed using an intention-to-treat approach, as used in the primary trials, where all data were included regardless of adherence to treatment. The intention-to-treat population included all randomized participants with an evaluable primary efficacy endpoint (baseline calculated LDL-C value and at least 1 post-baseline calculated LDL-C value up to Week 24). For most parameters, a mixed effects model with repeated measures (MMRM) was used to



(A)

Calculated LDL-C

(B)

ApoB

(C)

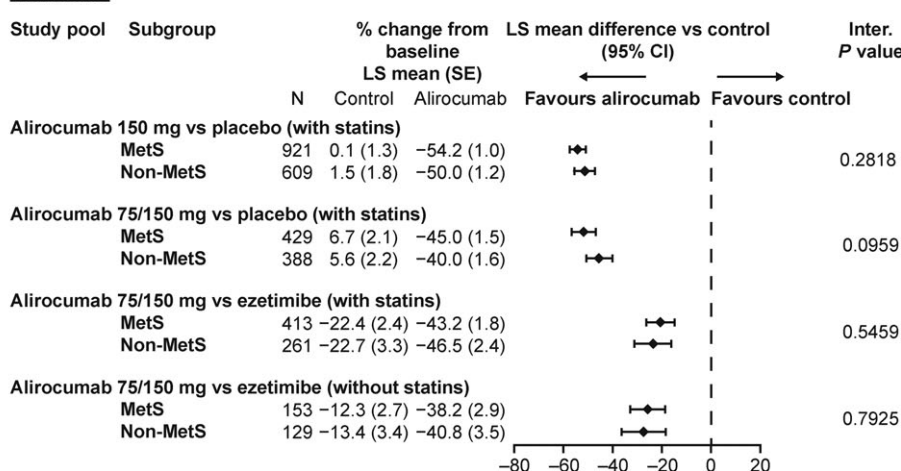
Non-HDL-C

FIGURE 2 (Continues)

account for missing data. Least squares means and standard errors (SE) were taken from the MMRM analysis. For analysis of TGs and Lp(a), adjusted means and SEs were taken from multiple imputation followed by robust regression. Interaction *P* values were calculated using the same model as above for comparing the difference (between alicumab and control) in percent change for lipid endpoints observed in both MetS and non-MetS patients. The proportion of patients achieving lipid goals was estimated from multiple imputation using only lipid data from patients who were on-treatment.

## 3 | RESULTS

## 3.1 | Patient characteristics

The randomized population for this analysis included 3341 patients (1940 with MetS; 1401 non-MetS); 1642 randomized individuals with diabetes were excluded. Baseline characteristics according to alicumab starting dose, control type and MetS status for each of the 4 pools are shown in Table 1 and online in Table S1. Patients had a mean age

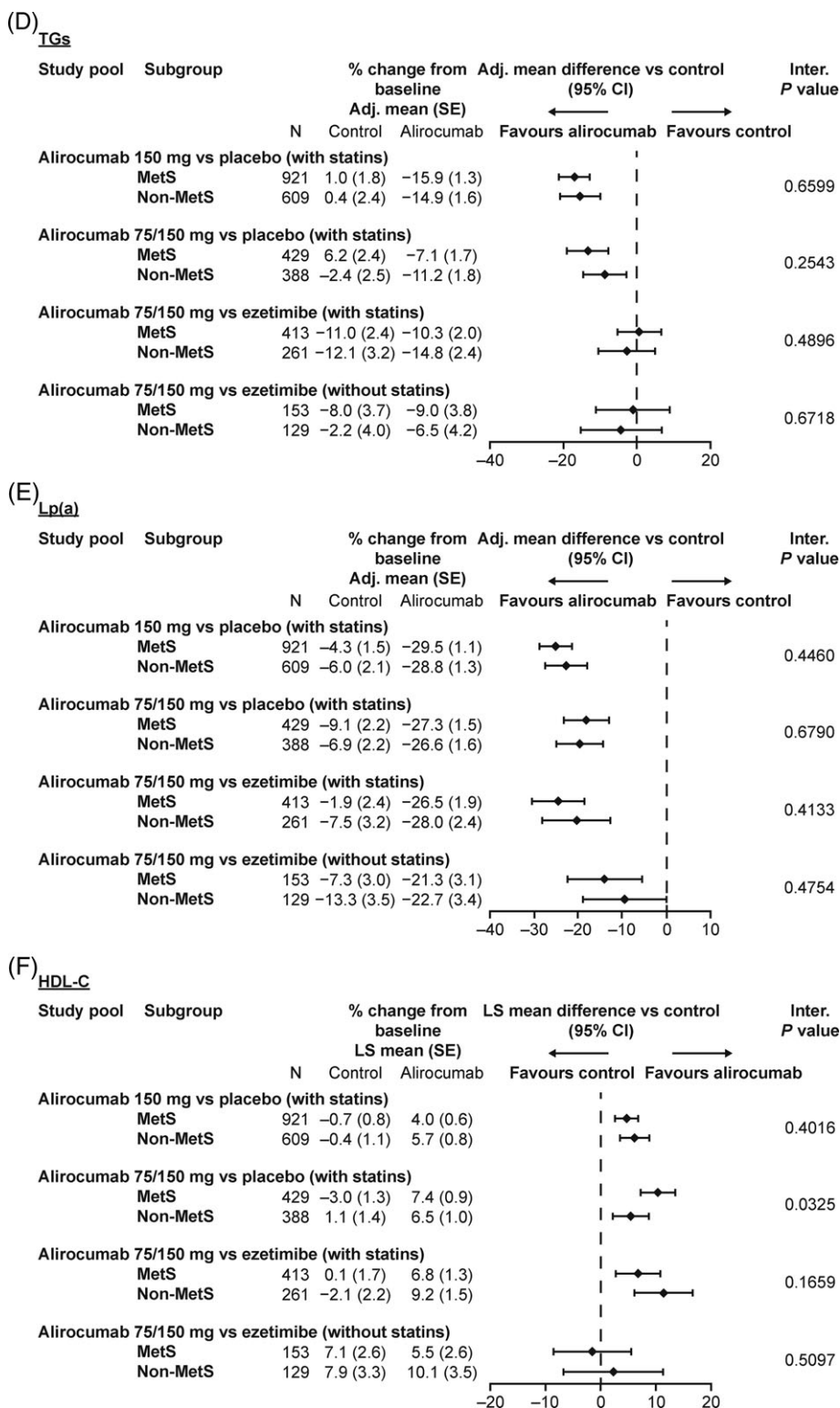


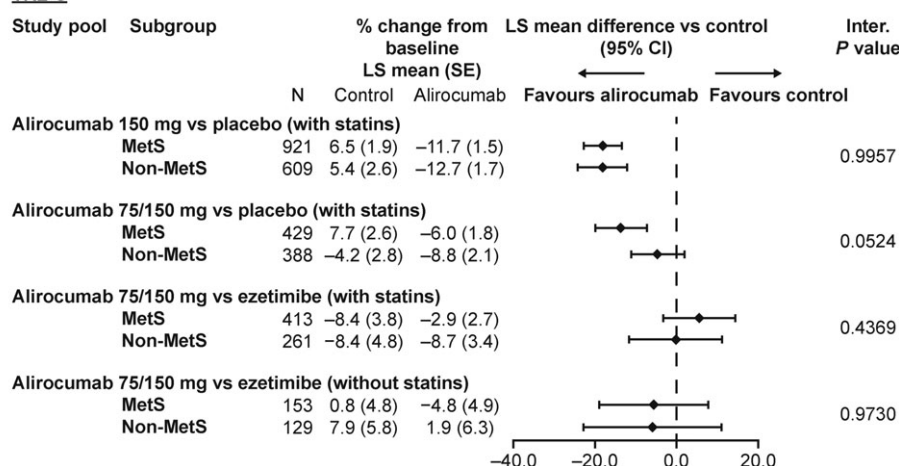
FIGURE 2 (Continues)

of 50 to 63 years across the groups; there was a greater proportion of males than females; and patients were mostly white (>85%). As per the selection criteria for this analysis, MetS patient groups had higher BMI, higher systolic blood pressure, a higher percentage of individuals with hypertension, higher FPG levels, lower levels of HDL-C, and higher levels of TGs compared to groups without MetS, for both placebo- and ezetimibe-controlled studies (Table 1). A higher proportion of patients with MetS were receiving other LLTs in addition to statin (Table S1).

Demographic and baseline characteristics were generally comparable between the alirocumab and control groups for each pool according to MetS status. However, for the ezetimibe-controlled studies, patients without MetS who received alirocumab in the 75/150 mg Q2W (with statins) pool were more likely to have hypertension (57.8% vs 46.0%) and atherosclerotic CV disease (90.4% vs 82.0%) than those in the control group; patients without MetS who received alirocumab in the 75/150 mg Q2W (without statins) pool were more likely to have

(G)

TRL-C



**FIGURE 2** Percent change from baseline in A, calculated LDL-C; B, ApoB; C, non-HDL-C; D, TGs; E, Lp(a); F, HDL-C and G, TRL-C at Week 24: Subgroup analysis according to MetS status at baseline (intention-to-treat analysis). Abbreviations: Apo, apolipoprotein; CI, confidence interval; HDL-C, high-density lipoprotein cholesterol; LDL-C, low-density lipoprotein cholesterol; Lp(a), lipoprotein(a); LS, least squares; MetS, metabolic syndrome; SE, standard error; TG, triglyceride; TRL-C, triglyceride-rich lipoprotein cholesterol. Non-HDL-C was calculated by subtracting HDL-C from total cholesterol. TRL-C was calculated by subtracting (HDL-C and calculated LDL-C) from total cholesterol

atherosclerotic CV disease (23.2% vs 15.7%) and less likely to be receiving other LLTs (10.0% vs 21.4%).

### 3.2 | Efficacy

LDL-C percentage reductions from baseline at Week 24 with alicumab were 63.9% (MetS) and 56.8% (non-MetS) in the pool of alicumab 150 mg Q2W (interaction  $P$  value  $<.05$ ), 52.2% (MetS) and 45.0% (non-MetS) in the pool of alicumab 75/150 mg Q2W vs placebo (interaction  $P$  value  $>.05$ ), 51.9% (MetS) and 52.6% (non-MetS) in the pool of alicumab 75/150 mg Q2W vs ezetimibe (with statins) (interaction  $P$  value  $>.05$ ), and 42.2% (MetS) and 46.7% (non-MetS) in the alicumab 75/150 mg Q2W pool (without statins) (interaction  $P$  value  $>.05$ ) (Figure 2A).

Across the 4 efficacy pools, by Week 24, alicumab reduced ApoB by 35.0% to 55.1% (MetS) and 35.9% to 51.3% (non-MetS) (Figure 2B), and non-HDL-C by 38.2% to 54.2% (MetS) and 40.0% to 50.0% (non-MetS) (Figure 2C). ApoB was reduced to a larger extent in the patients with MetS compared with non-MetS subjects in the alicumab 75/150 vs placebo pool only at Week 24 (interaction  $P$  values  $<.05$ ) (Figure 2B). For non-HDL-C, there was no difference in percent reduction from baseline at Week 24 for those with vs without MetS (Figure 2C). Apart from LDL-C percent reductions in the alicumab 150 mg Q2W pool (interaction  $P$  value  $<.05$ ), there were no differences in LDL-C, ApoB or non-HDL-C reductions between MetS and non-MetS groups at Week 12 (interaction  $P$  values  $>.05$ ) (Figure S1A-C).

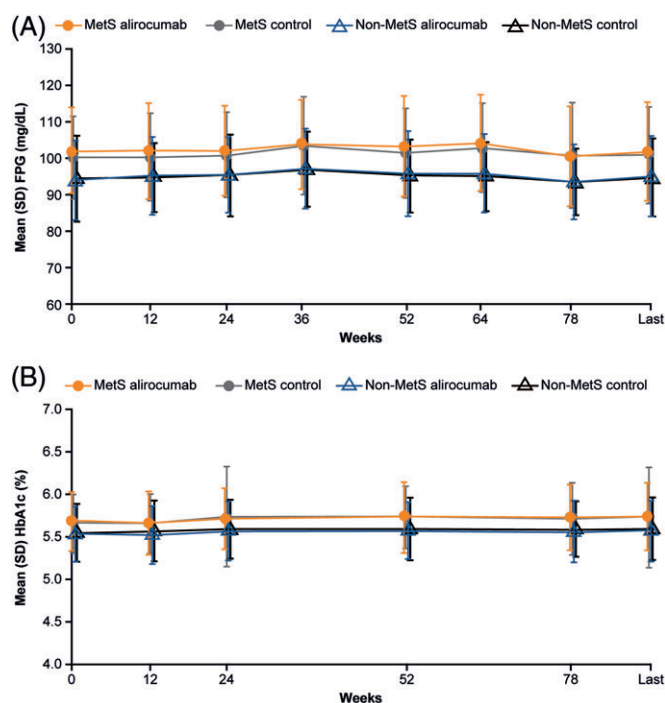
A greater proportion of patients receiving alicumab achieved LDL-C and non-HDL-C goals compared with controls at Week 24, regardless of MetS status (Tables S2 and S3).

Across the pools, by Week 24, TGs were reduced by 7.1% to 15.9% (MetS) and 6.5% to 14.9% (non-MetS) (Figure 2D), with similar results observed at Week 12 (Figure S1D). Lp(a) was reduced by 21.3% to 29.5% (MetS) and 22.7% to 28.8% (non-MetS) at Week 24 (Figure 2E). Reductions in each of these parameters were similar regardless of MetS status at Week 24 (interaction  $P$  values  $>.05$ ).

At Week 24, the use of other non-statin LLTs did not influence the percent change from baseline in TG levels observed with alicumab (Figure S2); for patients with MetS, the reduction from baseline was 13.3% to 22.6% (with LLT) vs 6.3% to 19.0% (without LLT) and for non-

MetS subjects, the reduction from baseline was 3.2% to 14.7% (with LLT) vs 0% to 10.0% (without LLT). Non-statin LLTs were examined in this analysis because of their potential effects on TG levels.

An increase in HDL-C levels of 4.0% to 10.1% was also observed across the 4 pools following alicumab therapy at Week 24 (Figure 2F). HDL-C was increased to a larger extent in the patients with MetS compared with the non-MetS subjects in the alicumab 75/150 mg vs placebo pool only (interaction  $P$  value  $<.05$ ); however, no significant difference between the MetS and non-MetS groups was observed at Week 12 (interaction  $P$  value  $>.05$ ) (Figure S1E). For



**FIGURE 3** Time profile of A, mean FPG and B, mean HbA1c levels: Subgroup analysis according to MetS status for pool of phase 3 studies (safety population). Abbreviations: FPG, fasting plasma glucose; HbA1c, glycated haemoglobin; MetS, metabolic syndrome; SD, standard deviation. Figures show combined pools for ezetimibe and placebo control pools, as differences between alicumab and control were similar in both pools. Last value defined as the last value collected up to 21 days after the last double-blind study treatment



patients with MetS or non-MetS subjects, the use of non-statin LLTs at Week 24 did not influence the percent change from baseline in HDL-C levels observed with alirocumab (Figure S3).

Across the 3 pools with patients on background statin therapy, alirocumab reduced TRL-C by 2.9% to 12.7% at Week 24, regardless of MetS status (Figure 2G). In contrast, for alirocumab-treated patients in the alirocumab 75/150 vs ezetimibe (without statins) pool, TRL-C was decreased (−4.8%) in patients with MetS and increased (+1.9%) in subjects without MetS (Figure 2G).

The percentage of patients for whom the alirocumab dose was increased from 75 to 150 mg Q2W at Week 12 was 36.1% (MetS) and 32.6% (non-MetS) in the 75/150 mg vs placebo pool, was 17.7% (MetS) and 15.1% (non-MetS) in the 75/150 mg vs ezetimibe with statin pool, and was 41.3% (MetS) and 35.0% (non-MetS) in the 75/150 mg vs ezetimibe without statin pool.

### 3.3 | Safety of alirocumab with respect to patients by MetS status

Time profiles of mean HbA1c and FPG levels over the treatment period showed that alirocumab had no clinically meaningful effect on these parameters compared to control treatments, for patients both with and without MetS (Figure 3). TEAE rates were generally similar across all treatment groups, irrespective of MetS status (Table 2). The most common TEAEs were nasopharyngitis, injection-site reaction, myalgia and upper respiratory tract infection. Injection-site reactions occurred more frequently in the alirocumab group vs the control group, for both placebo-controlled (6.5% vs 6.4% [MetS] and 11.1% vs 6.1% [non-MetS]) and ezetimibe-controlled (2.7% vs 2.0% [MetS] and 4.0% vs 1.8% [non-MetS]) trials (Table 2).

**TABLE 2** Safety analysis (safety population)

% (n)	Placebo-controlled studies				Ezetimibe-controlled studies			
	MetS (n = 1364)		Non-MetS (n = 1001)		MetS (n = 576)		Non-MetS (n = 395)	
	ALI (n = 893)	PBO (n = 471)	ALI (n = 687)	PBO (n = 314)	ALI (n = 329)	EZE (n = 247)	ALI (n = 226)	EZE (n = 169)
TEAEs	77.4 (691)	80.5 (379)	74.8 (514)	75.5 (237)	72.3 (238)	70.4 (174)	69.0 (156)	69.8 (118)
Treatment-emergent SAEs	13.7 (122)	13.6 (64)	11.8 (81)	9.9 (31)	12.2 (40)	12.6 (31)	13.7 (31)	8.9 (15)
TEAEs leading to death	0.3 (3)	0.4 (2)	0.7 (5)	1.3 (4)	0	0.8 (2)	0.4 (1)	1.8 (3)
TEAEs leading to discontinuations	4.5 (40)	5.3 (25)	4.9 (34)	4.1 (13)	9.7 (32)	8.1 (20)	8.0 (18)	11.2 (19)
TEAEs by preferred term in ≥5% individuals								
Nasopharyngitis	12.5 (112)	11.0 (52)	11.9 (82)	14.0 (44)	6.4 (21)	4.5 (11)	6.6 (15)	10.1 (17)
Injection-site reaction	6.5 (58)	6.4 (30)	11.1 (76)	6.1 (19)	2.7 (9)	2.0 (5)	4.0 (9)	1.8 (3)
Myalgia	5.5 (49)	3.8 (18)	4.7 (32)	4.8 (15)	8.5 (28)	8.9 (22)	6.6 (15)	7.7 (13)
Upper respiratory tract infection	5.4 (48)	6.2 (29)	6.3 (43)	7.0 (22)	7.3 (24)	6.9 (17)	4.9 (11)	3.6 (6)
Influenza	6.9 (62)	4.2 (20)	5.8 (40)	6.4 (20)	3.0 (10)	2.0 (5)	3.5 (8)	2.4 (4)
Headache	4.6 (41)	5.9 (28)	6.4 (44)	5.7 (18)	5.8 (19)	2.8 (7)	3.5 (8)	4.1 (7)
Arthralgia	5.3 (47)	5.9 (28)	4.2 (29)	5.7 (18)	4.6 (15)	4.9 (12)	4.9 (11)	3.0 (5)
Back pain	4.6 (41)	5.9 (28)	4.7 (32)	5.4 (17)	1.8 (6)	3.2 (8)	3.5 (8)	4.7 (8)
Diarrhoea	5.6 (50)	5.7 (27)	4.9 (34)	1.6 (5)	2.7 (9)	1.6 (4)	4.4 (10)	4.7 (8)
Dizziness	3.0 (27)	4.9 (23)	2.6 (18)	2.5 (8)	2.7 (9)	5.7 (14)	4.4 (10)	3.0 (5)
Urinary tract infection	4.8 (43)	3.8 (18)	4.8 (33)	5.1 (16)	2.1 (7)	4.0 (10)	0.9 (2)	2.4 (4)

Abbreviations: ALI, alirocumab; EZE, ezetimibe; PBO, placebo; MetS, metabolic syndrome; SAE, serious adverse event; TEAE, treatment-emergent adverse event.

## 4 | DISCUSSION

In this pooled analysis of 10 phase 3 ODYSSEY trials, alirocumab reduced levels of LDL-C, ApoB, Lp(a), TGs and non-HDL-C in patients with hypercholesterolaemia, regardless of MetS status. These findings are important because more than one-third of patients in the ODYSSEY trials had MetS (39%). This analysis demonstrated that the typical characteristics of these patients (mixed dyslipidaemia together with obesity, glucose intolerance and hypertension, most likely with additional concomitant medications) did not seem to have a major impact on the efficacy and safety of alirocumab. A slightly larger reduction in LDL-C was observed in individuals with MetS (vs no MetS) following alirocumab treatment in 1 of the study pools; however, as this was not observed in the other pools, it is unclear whether observed differences are real effects or the result of pooling the post-randomization groups with relatively small numbers of individuals.

The observed reductions in ApoB and non-HDL-C are of particular importance, as they may give a better estimate of the concentration of atherogenic particles than calculated LDL-C for patients with MetS because of the mixed dyslipidaemia profile.<sup>4,24</sup> It is interesting to note that alirocumab can increase the fractional clearance rates of non-LDL ApoB 100-containing particles such as intermediate density lipoprotein and, possibly, Lp(a).<sup>27</sup> The moderate reductions in TGs and moderate increases in HDL-C observed following alirocumab treatment also demonstrate an improvement in the lipid profile.

Patients with MetS had a higher rate of other LLT use, including increased fenofibrate, which is often prescribed to reduce TG levels.<sup>28</sup> However, use of non-statin LLTs in combination with alirocumab did not affect the moderate reductions in TGs seen with alirocumab treatment. In addition, the use of non-statin LLTs did not

affect the modest increases in HDL-C levels observed with alirocumab treatment.

Previous studies have shown that patients with MetS may not achieve their non-HDL-C goal following treatment with statins and other LLTs; in particular, the presence of MetS was shown to be a factor contributing to a greater difference between recommended vs attained non-HDL-C levels.<sup>8</sup> In the present analysis, a higher proportion of patients receiving alirocumab achieved both LDL-C and non-HDL-C goals compared with patients receiving placebo or ezetimibe, irrespective of MetS status.

With regards to safety, MetS status did not affect the incidence of TEAEs, with similar rates observed between the alirocumab and control groups. In this analysis, HbA1c and FPG levels were found to be unaffected by alirocumab in individuals both with and without MetS. Some studies have suggested that statin therapy is associated with an increased incidence of type 2 diabetes.<sup>29–31</sup> In addition, a Mendelian randomization study found that *PCSK9* and 3-hydroxy-3-methylglutaryl-coenzyme A reductase (*HMGCR*, the target of statin therapy) genetic variants associated with lower LDL-C were correlated with a reduced risk of CV events but also an increased risk of diabetes.<sup>32</sup> However, the authors emphasize that as *PCSK9* antibodies bind extracellular *PCSK9*, their biological effects may not be the same as those observed with *PCSK9* genetic variants which lower LDL-C levels over the course of a lifetime.<sup>32</sup> Furthermore, a pooled analysis of 10 ODYSSEY phase 3 studies, with follow-up periods of between 24 and 104 weeks, indicated that there was no evidence of an effect of alirocumab on the transition to new-onset diabetes in those individuals without diabetes or with pre-diabetes at baseline.<sup>33</sup> Also of note, subgroup analyses of ODYSSEY trials have shown no effect of diabetes on the efficacy of alirocumab in terms of LDL-C reduction from baseline up to Week 104.<sup>19,34</sup> Similarly, no effect of pre-diabetes on the efficacy and safety of alirocumab was observed.<sup>35</sup> Lastly, a prespecified analysis of the FOURIER clinical outcomes study with another *PCSK9* inhibitor, evolocumab, showed no effect of diabetic status on efficacy and safety with follow-up to 2.2 years, and no effect of evolocumab on the incidence of new-onset diabetes.<sup>36</sup>

Limitations of this analysis include the relatively short treatment periods of the ODYSSEY trials and the small number of patients in some subgroups. Waist circumference, a usual parameter for assessing obesity/MetS, was not measured in the trials. In addition, the analysis was performed on post-randomization groups, and some differences in baseline characteristics between the resulting alirocumab and control subgroups were noted. In order to provide sufficient numbers of patients for analysis, data were pooled from studies including patients with different clinical characteristics (eg, heterozygous familial hypercholesterolaemia,<sup>15,23</sup> non-familial hypercholesterolaemia with prior cardiovascular disease or other risk factors,<sup>17,18</sup> or statin intolerance<sup>22</sup>), which may limit interpretation of results.

In summary, this pooled analysis showed that alirocumab produced significant reductions in both LDL-C and non-HDL-C, of similar magnitudes in individuals both with and without MetS, and was generally well tolerated, with no apparent effect on measures of glycaemic control.

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## Conflict of interest

R. R. H. has received research funding from Abbott, AstaMed, Eli Lilly, Hitachi, Novo Nordisk, Sanofi-Lexicon and Viacyte; and has been a consultant/advisory panel member for Alere, AstraZeneca, Boehringer Ingelheim, Bristol Myers Squibb, Eisai, Elcelyx, Intarcia, Ionis, Janssen/Johnson & Johnson, Merck and Sanofi-Aventis. D. M.-W. has participated in speakers' bureaux for Amgen, Boehringer Ingelheim, Lilly, Sanofi, Merck (MSD), AstraZeneca and Novartis; and has been a consultant/advisory panel member for Sanofi, Boehringer Ingelheim, Merck, AstraZeneca and Novartis. P. R. T. has been a consultant/advisory panel member for Roche; and has participated in speaker's bureaux for Amgen, Regeneron Pharmaceuticals, Inc., Sanofi and Boehringer Ingelheim. M. B.-B. and A. L. are employees of/stockholders in Sanofi. M. J. L. is an employee of/stockholder in Regeneron Pharmaceuticals, Inc. H. N. G. has received research funding from Genzyme (Sanofi), Merck, Sanofi/Regeneron Pharmaceuticals, Inc. and Amgen; and has been a consultant/advisory panel member for Amarin, Amgen, AstraZeneca, Bristol Myers Squibb, GlaxoSmithKline, Ionis, Janssen, Kowa, Merck, Novartis, Sanofi/Regeneron Pharmaceuticals, Inc. and Pfizer.

## Author contributions

The sponsors were involved in the study design, collection, analysis and interpretation of data. R. R. H., D. M.-W., P. R. T., M. B.-B., M. J. L., A. L., and H. N. G. contributed to the data analysis and interpretation of the data, and critically reviewed and edited the manuscript. In addition, R. R. H., M. B.-B., M. J. L. and A. L. contributed to the concept or study design; and H. N. G. contributed to the data acquisition. All authors approved the final version. The authors had unrestricted access to study data, were responsible for all content and editorial decisions, and received no honoraria related to the development of this publication.

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## SUPPORTING INFORMATION

Additional Supporting Information may be found online in the supporting information tab for this article.

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